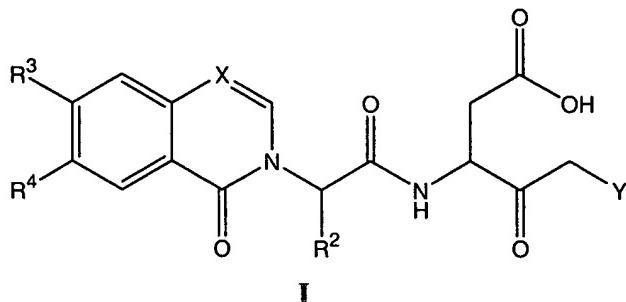


AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Previously presented) A compound of formula I:



X is CH;

Y is halo, trifluorophenoxy, or tetrafluorophenoxy;

R² is C₁₋₆ straight chained or branched alkyl;

R³ is hydrogen, halo, OCF₃, CN, or CF₃; and

R⁴ is hydrogen, halo, OCF₃, SR, CN, CF₃, Ar, or T-Ar; wherein:

T is O or S;

R is a C₁₋₆ straight chained or branched alkyl;

Ar is a phenyl ring optionally substituted with 1-3 groups selected from halo,

CH₃, CF₃, CN, OMe, OCF₃, and

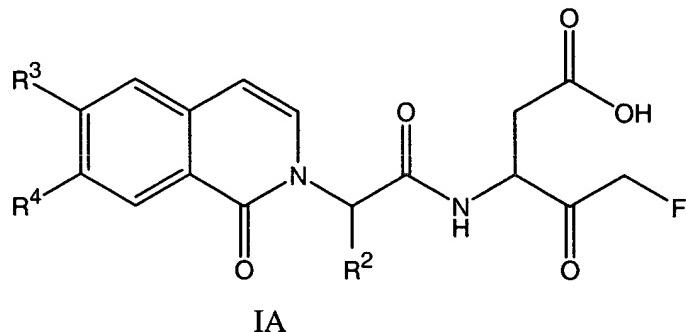
NR⁵R⁶; and

R⁵ and R⁶ each is independently H or C₁₋₆ straight chained or branched alkyl, or R⁵ and R⁶, taken together, form a 5-7 membered ring optionally containing up to 3 heteroatoms selected from O, S, NH, and N(C₁₋₆-straight chained or branched alkyl); provided that when Y is halo, then both, R³ and R⁴, are not simultaneously hydrogen.

2. (Original) The compound according to claim 1, wherein R² is ethyl, n-propyl, or isopropyl.

3. (Original) The compound according to claim 2, wherein Y is F, trifluorophenoxy, or tetrafluorophenoxy.

4. (Original) The compound according to claim 1, having formula IA:



wherein:

R² is ethyl, n-propyl, or isopropyl; and

R³ and R⁴ are each independently hydrogen, halo, OCF₃, CN, CF₃ or Ar, provided that both, R³ and R⁴, are not simultaneously hydrogen.

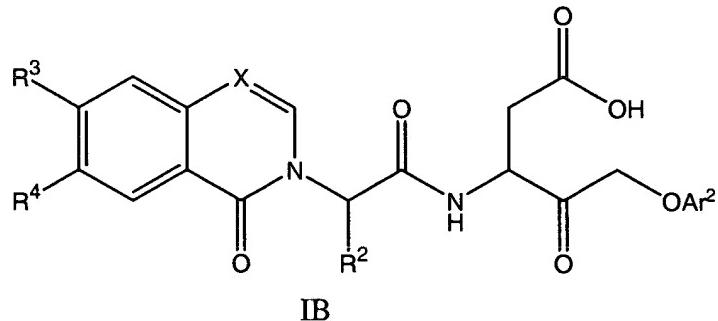
5. (Original) The compound according to claim 4, wherein R² is ethyl.

6. (Original) The compound according to claim 4, wherein R³ is hydrogen.

7. (Original) The compound according to claim 4 or claim 5, wherein R³ is H, and R⁴ is F, Cl, CN, Ar, or CF₃.

8. (Original) The compound according to claim 7, wherein R⁴ is Cl or CF₃.

9. (Previously presented) The compound according to claim 1, having the formula IB:



wherein:

X is CH;

R² is ethyl, n-propyl, or isopropyl;

R³ and R⁴ are each independently hydrogen, halo, OCF₃, CN, or CF₃; and Ar² is trifluorophenyl or tetrafluorophenyl.

10. (Original) The compound according to claim 9, wherein Ar² is 2,3,5,6-tetrafluorophenyl.

11. (Original) The compound according to claim 9, wherein R² is ethyl.

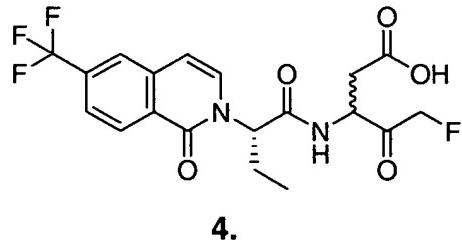
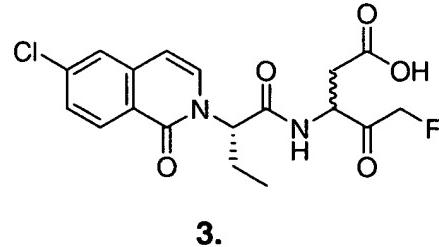
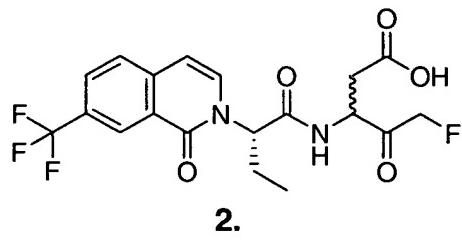
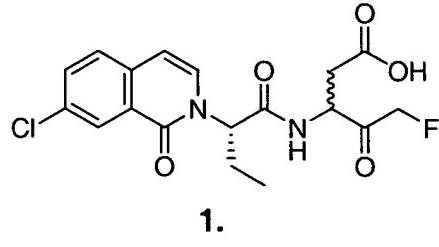
12. (Original) The compound according to claim 9, wherein X is CH.

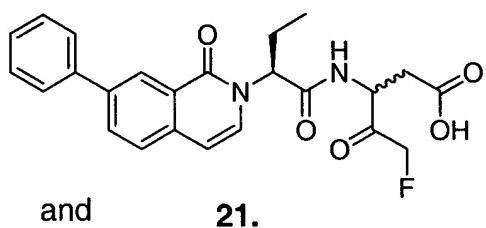
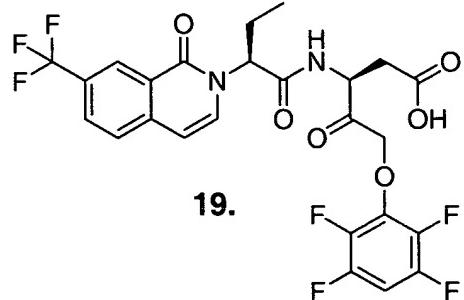
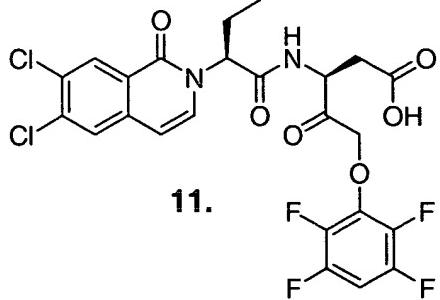
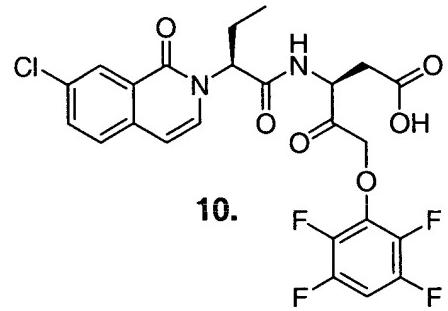
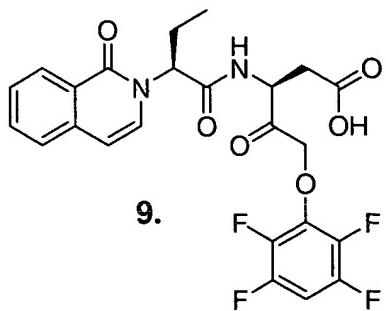
13. (Original) The compound according to claim 12, wherein R⁴ is Cl or CF₃.

14. (Original) The compound according to any one of claims 9-12, wherein R³ is H, and R⁴ is F, Cl, or CF₃.

15-19. (Canceled)

20. (Previously presented) The compound of claim 1, selected from:



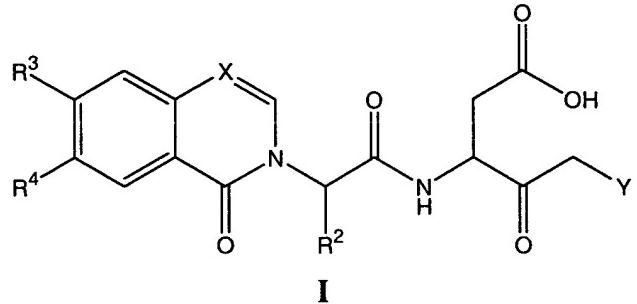


21. (Original) A pharmaceutical composition comprising:

- a compound according to claim 1; and
- a pharmaceutically acceptable carrier, adjuvant or vehicle.

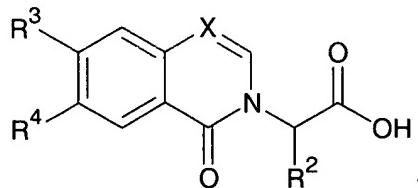
22-30. (Canceled)

31. (Previously presented) A method of preparing a compound of formula I,

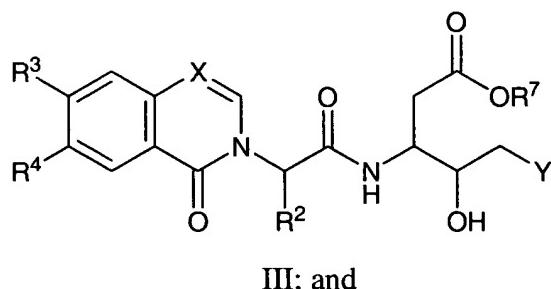
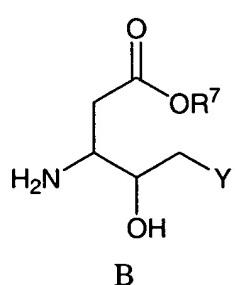


said method comprising:

reacting an acid or acid derivative of formula II,



with an amino alcohol of formula B, to provide a compound of formula III,



converting intermediate III to compound I, wherein;

X is CH;

Y is halo, trifluorophenoxy, or tetrafluorophenoxy;

R² is a C₁₋₆ straight chained or branched alkyl;

R³ is hydrogen, halo, OCF₃, CN, or CF₃; and

R⁴ is hydrogen, halo, OCF₃, SR, CN, CF₃, Ar, or T-Ar; wherein:

T is O or S;

R is a C₁₋₆ straight chained or branched alkyl;

Ar is a phenyl ring optionally substituted with 1-3 groups selected from halo,

CH₃, CF₃, CN, OMe, OCF₃, and

NR⁵R⁶;

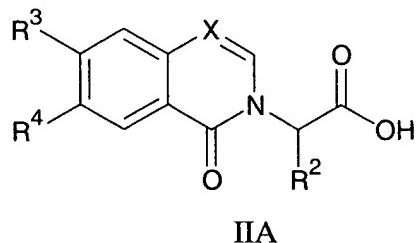
R⁵ and R⁶ each is independently H or C₁₋₆ straight chained or branched alkyl, or

R⁵ and R⁶, taken together, form a 5-7 membered ring optionally containing up to 3 heteroatoms selected from O, S, NH, and N(C₁₋₆ straight chained or branched alkyl);
and

R⁷ is a suitable protecting group;

provided that when Y is halo, then both, R³ and R⁴, are not simultaneously hydrogen.

32. (Previously presented) A compound of formula IIA:



wherein;

X is CH;

R² is a C₁₋₆ straight chained or branched alkyl;

R³ is hydrogen, halo, OCF₃, CN, or CF₃; and

R⁴ is hydrogen, halo, OCF₃, SR, CN, CF₃, Ar, or T-Ar; wherein:

T is O or S;

R is a C₁₋₆ straight chained or branched alkyl;

Ar is a phenyl ring optionally substituted with 1-3 groups selected from halo, CH₃, CF₃, CN, OMe, OCF₃, and

NR⁵R⁶; and

R⁵ and R⁶ each is independently H or C₁₋₆ straight chained or branched alkyl, or R⁵ and R⁶, taken together, form a 5-7 membered ring optionally containing up to 3 heteroatoms selected from O, S, NH, and N(C₁₋₆-straight chained or branched alkyl).

33. (Currently amended) The compound according to claim 31 or 32 wherein R² is ethyl or isopropyl.